

K024170 L1 SYSTEMApr 14, 2003
117 days to decisionK024170 · Product code: **FMK** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k024170/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Single Use Only Blood Lancet With An Integral Sharps Injury Prevention Feature (FMK)
Date received	Dec 18, 2002
Decision date	Apr 14, 2003
Days to decision	117 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Pelikan Technologies, Inc.
Location	Palo Alto, CA, US
Contact	JACK GREEN
510(k) history	2 submissions · 2 cleared · 2003-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k024170/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026